IN THE SPECIFICATION

Please delete the paragraph inserted by the Preliminary Amendment of August 30, 2001 at page 1, line 4, and insert the following paragraph:

This application is a division of U.S. application Serial No. 08/945,289, filed October 17, 1997, in issue, which is a continuation-in-part of U.S. application Serial No. 08/432,483, filed May 1, 1995, now U.S. Patent 6,410,022.

A version of the amended paragraph with markings on a separate sheet pursuant to 37 C.F.R. § 1.121(b) is provided as Exhibit A.

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IN THE CLAIMS

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Please amend the claims to appear as written below pursuant to 37 C.F.R. § ECH CENTER 1600/2900 1.121(c)(1)(i). A version of the amended claims with markings on a separate sheet pursuant to 37 C.F.R. § 1.121(c)(1)(ii) is provided as Exhibit B, and a complete set of pending claims, as amended herein, is provided pursuant to 37 C.F.R. § 1.121(c)(3) as Exhibit C.

Amended Claims

- 28. (amended) A method for treating or preventing atherosclerosis in a human or animal comprising administering to said human or animal an antigenic vaccine peptide comprising a universal helper T cell epitope portion linked to a B cell epitope portion, wherein said B cell epitope portion comprises a B cell epitope of CETP.
- 29. (amended) The method according to claim 28, wherein said helper T cell epitope portion comprises a helper T cell epitope derived from an antigenic peptide selected from the group consisting of tetanus toxoid, diphtheria toxoid, pertussis vaccine, Bacile Calmette-Guerin (BCG), polio vaccine, measles vaccine, mumps vaccine, rubella vaccine, purified protein derivative of tuberculin, keyhole limpet hemocyanin, hsp70, and combinations thereof.
- 37. (amended) The method according to claim 28, wherein said B cell epitope portion of the antigenic vaccine peptide comprises 6 to 26 consecutive amino acids of the carboxyl terminal 26 amino acids of human cholesteryl ester transfer protein (SEQ ID NO:1).